

AMENDMENTS TO THE CLAIMS

1. (Currently amended) A pharmaceutical composition suitable for oral administration in the form of a homogeneous solution which on exposure to water or gastrointestinal fluids forms an emulsion having a particle size of less than 5 microns, the solution comprising:

~~(b)~~ (a) a pharmaceutically effective amount of a cyclosporin,

~~(e)~~ (b) a carrier medium comprising a triglycerol monoester of a fatty acid having from 6 to 30 carbon atoms or mixtures thereof,

~~(d)~~ (c) polyethylene glycol,

~~(e)~~ (d) a non-ionic surfactant having a hydrophilic lipophilic balance (HLB) greater than 10, and

~~(f)~~ (e) optionally, a viscosity reducing agent,

the composition being substantially free from ethanol.

2. (Original) A pharmaceutical composition according to claim 1, wherein said fatty acid has 8 to 18 carbon atoms.

3. (Original) A pharmaceutical composition according to claim 1, wherein the carrier medium comprises a triglycerol monoester of capric acid, caprylic acid, lauric acid, oleic acid, or a mixture thereof.

4. (Original) A pharmaceutical composition according to claim 1, wherein the carrier medium comprises triglycerol monooleate.

5. (Original) A pharmaceutical composition according to claim 1, wherein the cyclosporin is 1 to 25% by weight of the composition, the carrier medium is 15-50% by weight of the composition, the non-ionic surfactant is 5-40% by weight of the composition, the polyethylene glycol is 5 to 40% by weight of the composition, and the viscosity reducing agent, when present, is 5 to 25% by weight of the composition.
6. (Original) A pharmaceutical composition according to claim 5 wherein the cyclosporin is 5 to 20% by weight of the composition, the carrier medium is 20-40% by weight of the composition, the non-ionic surfactant is 10-30% by weight of the composition, and the polyethylene glycol is 10 to 35% by weight of the composition.
7. (Original) A pharmaceutical composition according to claim 6, wherein the cyclosporin is 10 to 20% by weight of the composition, the carrier medium is 25-35% by weight of the composition, the non-ionic surfactant is 15-25% by weight of the composition, the polyethylene glycol is 20 to 30% by weight of the composition, and the viscosity reducing agent, when present, is 10 to 20% by weight of the composition.
8. (Original) A pharmaceutical composition according to claim 5, wherein the amount of cyclosporin is 5 to 400 mg, said fatty acid has 8 to 18 carbon atoms, the non-ionic surfactant has a HLB greater than 12, and the polyethylene glycol has a molecular weight of 200 to 1,000.
9. (Original) A pharmaceutical composition according to claim 8, wherein the amount of cyclosporin is 20 to 200 mg, the non-ionic

surfactant has a HLB greater than 14, and the polyethylene glycol has a molecular weight of 200 to 600.

10. (Original) A pharmaceutical composition according to claim 8, wherein the non-ionic surfactant makes up from 15 to 25% by weight of the composition.

11. (Original) A composition according to claim 1, wherein the polyethylene glycol has a molecular weight of from 200 to 1000.

12. (Original) A pharmaceutical composition according to claim 1, wherein the non-ionic surfactant is selected from the group consisting of: polyoxyethylated hydrogenated vegetable oils, polyethoxylated castor oils, polyethoxylated hydrogenated castor oil, polyoxyethylene-sorbitan-fatty acid esters, and polyoxyethylene castor oil derivatives.

13. (Original) A pharmaceutical composition according to claim 12, wherein the non-ionic surfactant is selected from the group consisting of polyoxyethylene (20) sorbitan monolaurate, polyoxyethylene (20) sorbitan monopalmitate, polyoxyethylene (20) sorbitan monostearate, polyoxyethylene (20) sorbitan monooleate, PEG-30 hydrogenated castor oil, PEG-40 hydrogenated castor oil, PEG-50 hydrogenated castor oil, PEG-60 hydrogenated castor oil, polyoxyethylene 40 castor oil, polyoxyethylene 60 castor oil, polyoxyethylene 35 castor oil, and mixtures thereof.

14. (Original) A pharmaceutical composition according to claim 1, wherein the viscosity reducing agent is present and is selected from the group consisting of monoesters of glycerol and aliphatic monocarboxylic acids having from 6 to 30 carbon atoms, and mixtures thereof.

15. (Currently amended) A pharmaceutical composition according to claim ~~15~~ 14, wherein a viscosity reducing agent is present and selected from the group consisting of glycerol monocaprylate, glycerol monooleate, and mixtures thereof.

16. (Original) A pharmaceutical composition according to claim 1, further comprising an antioxidant.

17. (Original) A composition according to claim 1, wherein the weight ratio of the carrier medium, non-ionic surfactant and viscosity reducing agent taken together to polyethylene glycol is greater than 1.

18. (Currently amended) A pharmaceutical composition according to claim ~~18~~ 17, wherein the cyclosporin is 1 to 25% by weight of the composition; the carrier medium is 15-50% by weight of the composition and comprises a triglycerol monoester of capric acid, caprylic acid, lauric acid, oleic acid, or a mixture thereof; the non-ionic surfactant is 5-40% by weight of the composition and is selected from the group consisting of: polyoxyethylated hydrogenated vegetable oils, polyethoxylated castor oils, polyethoxylated hydrogenated castor oil, polyoxyethylene-sorbitan-fatty acid esters, and polyoxyethylene castor oil derivatives; the

polyethylene glycol is 5 to 40% by weight of the composition and has a molecular weight of 20 to 1,000; the viscosity reducing agent, when present, is 5 to 25% by weight of the composition and is selected from the group consisting of monoesters of glycerol and aliphatic monocarboxylic acids having from 6 to 30 carbon atoms, and mixtures thereof; and optionally, an antioxidant is in an amount of from 0.01% to 2% by weight of the total composition and is being selected from the group consisting of BHA, BHT, and alpha-tocopherol.

19. (Original) A pharmaceutical composition according to claim 11 wherein the cyclosporin is Cyclosporin A.

20. (Original) A pharmaceutical composition according to claim 1, formulated as a drinking solution.

21. (Original) A pharmaceutical composition according to claim 1, formulated as a hard or soft capsule.

22. (Original) A pharmaceutical composition according to claim 1, contained within a soft gelatine capsule.